Kazuo Suzuki*: Biosystematic studies of Japanese Epimedium (Berberidaceae)

(1) Variation of the populations in Shikoku (Part 2)**

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Populations showing problematical affinity and/or variability. It has been confirmed that eight of the 23 populations studied can not be adequately referred to any one of the four representative types described above. In the eight populations were included the individuals which are deviated from "pure" species in the combination of morphological characteristics, or are intermediate in some features. The individuals, which can be identified to belong to one of the four species, were also mixed in most of the eight populations.

In the following descriptions which outline the population structure, individuals referable to "pure" species are represented by the initial of species name (e.g., *E. diphyllum*=D), and the individuals which show the morphological traits being laid across two species are denoted by the two initials connected by a hyphen. The number of samples observed is inserted in parentheses.

Koboshi (Fig. 8, 2): varying from D to K, also skewing to T:D (7); K (1); D-K (9), D in leaf hairiness and K in ramification and/or leaflet apex; nearly T (1), differing from typical T in having a shorter spur; D-T (2), D in ramification and T in spur development (though shorter than typical T) and leaflet apex.

Masahiro (Fig. 8, 3): varying from D to K: D(2); K(4); D-K(6), D in leaf hairiness and K in ramification and/or leaflet apex.

Tōzenji (Fig. 8, 5): close to K but faintly skewing to D: K(14); K-D(10), D in leaflet apex and K in other features.

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^{**} Continued from Journ, Jap. Bot. 53: 203-212 (1978).

Kagami-A (Fig. 8, 9): close to D but faintly skewing to T:D(3), D-T(7), T in leaflet apex and D in other features.

Kagami-B (Fig. 8, 10): intermediate between D and T though closer to D, variable: D (5); D-T (25), D in ramification and T in spur development (though generally shorter than typical T) and/or leaflet apex.

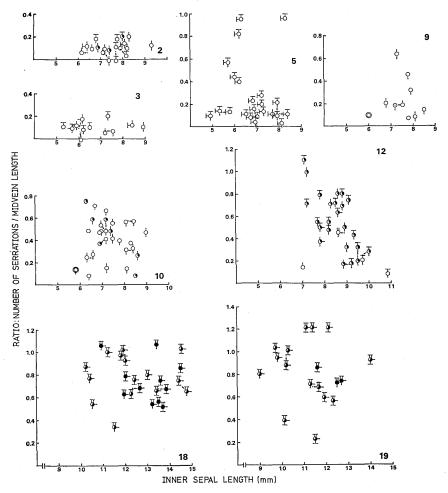


Fig. 8. Scatter diagrams showing the character correlation in the collections from eight problematical populations: 2, Koboshi; 3, Masahiro; 5, Tôzenji; 9, Kagami-A; 10, Kagami-B; 12, Aburaishi; 18, Sadanaga; 19, Funakoshi. (Legends for symbols as in Fig. 5).

Aburaishi (Fig. 8, 12): intermediate between D and T though closer to T, variable: D(1); D-(T)-G(1), D in spur development, G in ramification and T (intermediate between D and G) in other features; D-T (2), D in spur development, and T in ramification and leaflet apex; nearly T (24), differing from typical T in having a shorter spur.

A herbarium specimen from Ioki (cf. Fig. 1 and Fig. 4); D-T, D in spur development and T in other features.

A herbarium specimen without a flower from Kuwao (cf. Fig. 1 and Fig. 4); T-G, T in leaflet apex, intermediate in leaf hairiness, and G in ramifica-

Table 3. Number of individuals showing respective pollen fertility.

Locality		Pollen fertility (%)						
		40-50	50-60	60-70	70-80	80-90	90-100	undeterminable
1.	Mt. Iino	0	0	0	0	0	5	0
2.	Koboshi	0	0	0	1	0	19	0
3.	Masahiro	0	0	1	1	0	10	0
4.	Kirihata	0	0	0	0	1	28	0
5.	Tōzenji	0	0	0 -	1	0	23	0
6.	Nishikata	0	0	0	2	4	14	0
7.	Takano	0	0	0	0	0	8	0
8.	Go-ōdō	0	1	1	0	0	15	1
9.	Kagami-A	0	0	0	0	0	10	1
10.	Kagami-B	0	0	0	0	5	25	0
11.	Sako	0	0	0	0	1	17	0
12.	Aburaishi	0	0	0	2	2	24	0
13.	Okō	0	0	0	0	0	26	0
14.	Ikku	0	0	0	0	0	21	0
15.	Jinzenji	0	0	0	0	0	10	0
16.	Engyōji	0	0	0	0	0	27	0
17.	Fukui	0	1	0	0	0	22	0
18.	Sadanaga	0	0	0	1	0	25	0
19.	Funakoshi	0	0	0	0	0	20	0
20.	$ar{\mathrm{O}}$ nogahara	1	0	0	0	2	20	. 1
21.	Yamada	0	0	0	0	0	27	2
22.	Izumi	0	0	0	2	1	10	11
23.	Misaki	0	0	0	1	0	34	0

tion.

Sadanaga (Fig. 8, 18): intermediate between T and G, showing no individual of typical T or G but composed of the individuals having the following character-combinations (T and G are denoted below as the features of the corresponding taxon, and I means the intermediate trait between T and G; the characters are arranged in the order of the degree of spur development, leaf hairiness, type of ramification and shape of a leaflet apex): TIGG (11), TIGT (5), GIGG (4), GIGT (3), GTGT (1), TTGG (1).

Funakoshi (Fig. 8, 19): intermediate between T and G: no typical T nor G; composed of the individuals with the following character combinations (procedures as in the Sadanaga population): TIGG (12), TIGT (3), GIGT (2), GIGG (1).

Pollen fertility. Results of the examination of pollen are shown in Table 3. It is clear that most individuals in every population show over 80% in pollen fertility and that individuals with an aberrant character-combination in problematical populations show as high fertility as those in the populations of the representative types.

Discussion The present work has proved that populations of *Epimedium* in Shikoku can be divided into two kinds. One of them has a minor intrapopulational variability and consists of the populations which can be identified as belonging to one of the four representative types. The other kind of the populations is characterized by a larger intra-populational variability. They are composed of the individuals which are referable to a representative type and some aberrant individuals or of aberrant individuals only.

Character-expressions found in such aberrant individuals strongly suggest that gene exchanges among the four types would have occurred. It is hardly imagined that the intra-populational variability as noted in the Sadanaga population has been brought about through mutation or phenotypic plasticity. More or less similar variability within a population, which can be regarded as a reproductive unit (cf. Table 1), has been observed in seven other populations. It may be presumed that the gene-recombinations through hybridization have occurred.

As may be seen from Table 2 and Fig. 5, *E. diphyllum* and *E. grandi*florum are contrasted in all the diagnostic characters, except the hairiness of the upper leaf surface (see below). The morphological features of these two species can be regarded to represent the two extremes among plants of *Epimedium* in Shikoku. On the other hand, *E. kitamuranum* and *E. trifoliato-binatum* are intermediate between the two extremes in various features and approach to one of the two extremes in some other features. It may also be clear from Table 2 that *E. kitamuranum* is closer to *E. diphyllum* while *E. trifoliatobinatum* to *E. grandiflorum*. In case of *Epimedium*, sterility barriers apparently do not operate to reduce introgressive hybridization, as evidenced by the fact that no appreciable reduction pollen- (Table 3) and seed-fertility has been detected in the aberrant individuals of various variable populations. All of these observations are in favor of the hypothesis advanced by some earlier authors (Stearn, 1938; Maekawa, 1955; Kitamura & Murata, 1961) that both *E. kitamuranum* and *E. trifoliatobinatum* may be of hybrid origin between *E. diphyllum* and *E. grandiflorum*.

Mention should be made to the hairy upper surface of a leaf in E. kitamuranum. This trait is not found in any other plants of Epimedium in Shikoku, and it would seemingly contradict the assumption of the hybrid origin of E. kitamuranum. The hairs in question are, however, morphologic-

ally similar to the hairs on the lower surface of a leaf in *E. kitamuranum* and also in *E. diphyllum* and *E. trifoliatobinatum* (Fig. 9). Morphological characteristics of hybrids are not always parental or intermediate, but transgressive segregation is known in various examples (Grant, 1975). The fact that only *E. kitamuranum* has the hairy upper leaf surface among the plants in Shikoku may therefore be not necessarily contradictory to the assumption of its hybrid origin.

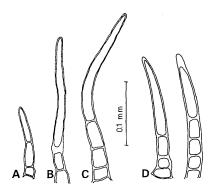


Fig. 9. Persistent hairs observed on a leaf: A, E. diphyllum; B, E. kitamuranum; C, E. trifoliatobinatum; D, E. kitamuranum. A-C, lower surface; D, upper surface.

Although the above hypothesis seems to be tenable, it is apparently desirable to verify it by experimental means. It must be noted that no population which is indicative of natural hybridization between $E.\ diphyllum$ and $E.\ grandiflorum$ has been located in Shikoku. Under the assumption that E.

kitamuranum and E. trifoliatobinatum are hybrid derivatives, studies must be extended to make clearer what environmental factors have contributed to sort out the plants referable to E. kitamuranum and E. trifoliatobinatum from the hybridizing populations. Edaphic factors may be regarded as part of such factors; "pure" populations of E. trifoliatobinatum are almost always confined to a site on serpentine rocks according to Yamanaka (1953a) and the present observation.

Evolutionary differentiation which seems not to be related with hybridization has been observed in *E. grandiflorum*. Populations of this species in eastern Shikoku (7 and 8) are characterized by a very short petiole length of a stem-leaf, while a population located in western Shikoku (20) has consisted of the individuals with much longer petiole. The former grows on limestone cliffs or slopes, while the latter occurs on a grassy karst plateau. These differentiation must have occurred in connection with geographical isolation, and hybridization seems not to have played a role. This subject, as well as various other subjects, should be studied further by adding analyses of populations in other regions and by approaches from other points of view. Studies along these lines are now in progress.

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四国産イカリソウ属植物の23集団からサンプリングを行い、形態的変異及び花粉の 稔性について調べた。その結果、現在見られる様々な集団の構成パターンの成立に浸 透性交雑が関与していたものと思われる。サイコクイカリソウ、ヒメイカリソウと呼 ばれる種類はバイカイカリソウとヤチマタイカリソウ(広義)との間の自然交雑を起源 とし現在は比較的安定していると思われる。一方、交雑が関与していないと思われる2 つの変異型が地理的に隔離されたヤチマタイカリソウの集団間に認められる。